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CLAIMS

- 1. A process for purification of a heterologous protein of interest, comprising
- 5 (a) providing a fusion protein comprising said heterologous protein fused to a CBM intercepted by a proteolytic cleavage site,
 - (b) contacting said fusion protein with a functional protease fused to a CBM , at conditions facilitating proteolytic cleavage by said protease, to cleave the CBM from the heterologous protein of interest,
- (c) contacting the solution of CBM-protease, free CBM and heterologous protein of interest to a polysaccharide matrix, under conditions where the CBMprotease and free CBM binds to said polysaccharide matrix and where the heterologous protein of interest is not retained on said polysaccharide matrix,
 - (d) separating the non-bound heterologous protein of interest from the polysaccharide matrix,
 - (e) washing the polysaccharide matrix with the bound CBM-protease and CBM, with one or more suitable aqueous solutions,
 - (f) eluting the CBM-protease from the matrix by adjusting conditions effecting the release of said CBM-protease off the matrix; and
- (g) optionally reconditioning said eluted CBM-protease, to retain its affinity to said polysaccharide matrix, such that the reconditioned CBM-protease can be re-used for subsequent repetition of the process defined by steps (a) (g) wherein said CBMs are capable of binding reversibly to a polysaccharide matrix and being released from such matrix by non-denaturing elution conditions and do not bind substantially to insoluble cell-wall plant material, the method.
 - 2. The process of claim 1, wherein said protease fused to CBM is from the group of proteases cconsisting of enterokinase, tobacco etch virus (TEV) protease, factor X and thrombin.
 - 3. The process of claim 2 wherein said protease is mammalian enterokinase (EK) or an enterokinase active part thereof.
- 4. The process of claim 3, wherein said EK comprises a bovine EK catalytic domain (EKc).
 - The process of claim 4, wherein said bovine EKc is encoded by the nucleic acid sequence shown as SEQ ID NO: 2.